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(54) Title: ONE-PART ORGANOPOLYSILOXANE RUBBER COMPOSITION FOR USE AS A CORROSION PROTECTION COATING ON METALS

(57) Abstract

The present invention relates to a one-part room temperature vulcanizable organopolysiloxane rubber composition which crosslinks in the presence of moisture to form a coating for the corrosion protection of metals. The one-part organopolysiloxane rubber composition comprises the product which is obtained by mixing the following: a) about 20 to 50 weight percent of polydimethyl siloxane fluid of the formula: HO[(CH₃)₂SiO]_nH, in which n has an average value such that the viscosity is in the range from 1 to 100 Pa.S at 25 °C, preferably from 10 to 20 Pa.S at 25 °C; b) about 20 to 50 weight percent of a mixture of anorphorous and crystalline SiO₂ reinforcing filters having a surface area of up to 200 m²/g and a specific gravity of 2.2; c) about 2 to 6 weight percent of an oximino silane; d) about 0.25 to 2 weight percent of an organo functional silane; e) about 0.05 to 1 weight percent of an organotin salt; and f) about 10 to 40 weight percent of an organic solvent as a dispersion medium for the above composition.

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TITLE: ONE-PART ORGANOPOLYSILOXANE RUBBER COMPOSITION
FOR USE AS A CORROSION PROTECTION COATING ON METALS

FIELD OF THE INVENTION

This invention relates to a one-part room temperature vulcanizable organopolysiloxane rubber composition which crosslinks in the presence of moisture to form a coating for the corrosion protection of metals.

10 BACKGROUND OF THE INVENTION

Metal surfaces exposed to moisture such as rain or fog in combination with contaminated atmospheres as are found in industrial locations may be subject to extensive corrosion unless protected in some way from exposure to the corrosive atmosphere. Other potentially corrosive 15 environments include along sea coasts where salt spray is found and in areas where agricultural chemicals are widely distributed. In addition, metal surfaces directly exposed to water such as marine structures and vessels are also subject to the potential for extensive corrosion. 20 past, such metal surfaces have been most commonly protected by being painted with alkyd based paints. Such paints form a relatively rigid coating on the surface of the metal which can become brittle and when subjected to stress, can flake or chip off, thereby exposing the underlying metal to 25 the corrosive elements. In addition, such paints generally are susceptible to UV damage thereby further reducing their effective life.

Two-part organopolysiloxane rubber compositions

for use as a corrosion protection coating on metals have
been developed. For example, Lampe describes in U.S.

Patent No. 4,341,842 a two-part room temperature
vulcanizable composition for coating the underside of

vehicles to protect the metal from rusting or being corroded by road salts or other similar compounds. However, such two-part compositions have a major disadvantage in that they require the use of complex dual mixing and spray nozzle apparatus or require pre-mixing and immediate use on site when used with conventional spray equipment. If conventional spray equipment is used, the amount of material pre-mixed must also be exact to prevent wastage as the composition has a finite pot life.

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SUMMARY OF THE INVENTION

In one aspect the present invention provides for a one-part room temperature vulcanizing organopolysiloxane rubber composition for use as a corrosion protection coating on metals. The composition comprises the product which is obtained by mixing the following:

> a) about 20 to 50 weight percent of polydimethyl siloxane fluid of the formula

$HO[(CH_3)_2SiO]_nH$

- in which n has an average value such that the viscosity is in the range from 1 to 100 Pa.S at 25°C, preferably from 10 to 20 Pa.S at 25°C;
 - b) about 20 to 50 weight percent of a mixture of anorphorous and crystalline SiO₂ reinforcing fillers having a surface area of up to 200 m²/g and a specific gravity of 2.2;
 - about 2 to 6 weight percent of an oximino silane;
 - d) about 0.25 to 2 weight percent of an organo functional silane;
- 30 e) about 0.05to 1 weight percent of an organotin salt; and

f) about 10 to 40 weight percent of an organic solvent as a dispersion medium for the above composition.

The present invention also provides for a method of protecting exposed surfaces particularly metal or concrete surfaces from the effects of a corrosive environment. The method comprises applying to the surface a thin layer of the above one-part organopolysiloxane rubber composition and allowing the layer of the one-part organopolysiloxane rubber composition to cure at room temperature to a silicone elastomer.

The present invention also provides for the metal or concrete surfaces coated with the silicone elastomer formed from the curing of the one-part organopolysiloxane rubber composition.

DETAILED DESCRIPTION OF THE INVENTION

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The one-part organopolysiloxane rubber compositions of the present invention are ideally suited for corrosion protection of steel or concrete structures against salt spray and chemical environments including direct exposure to salt water, salt fog and other industrial pollutant. The compositions of the present invention can be used to coat metal surfaces of motor vehicles which may be exposed to high salt condition during the winter season. The compositions of the present invention are particularly useful on marine installations, electric transmission towers and bridges for corrosion protection of steel or concrete structures directly exposed to salt water and industrial pollution, especially sulfur based.

The one-part organopolysiloxane rubber compositions of the present invention for use as a corrosion protective coating on metals contain about 20 to

50 weight percent of polydimethylsiloxane fluid of the formula $HO[(CH_3)_2SiO]_nH$ in which n has an average value such that the viscosity is in the range from 1 to 100 Pa.S, preferably 10 to 80 Pa.S, most preferably 10 to 50 Pa.S at $25^{\circ}C$.

The composition also contains about 20 to 50 weight percent of a mixture of anorphorous and crystalline SiO_2 reinforcing fillers having a surface area of up to 200 m²/g and preferably a specific gravity of 2.2.

The composition also contains about 2 to 6 weight percent of an oximinosilane cross linking agent.

Preferably the oximinosilane cross linking agent is of the formula RSi(ON=CR'2)3 in which R and R' each represent a monovalent hydrocarbon radical such as for example an alkyl radical such as methyl, ethyl, propyl, butyl, or an alkylene radical such as vinyl and allyl. The preferred R and R' are alkyl radicals, most preferably methyl and ethyl radicals.

The composition also contains about 0.25 to 2

weight percent of an organo functional silane. Preferably the organo functional silane has the formula

$$R^{3}_{b}$$
 | | 25 | $(R^{2}0)_{3-b} - Si - Z$

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wherein R^2 and R^3 are C_{1-8} monovalent hydrocarbon radicals, b varies from 0 to 3, preferably 0, Z is a saturated, unsaturated or aromatic hydrocarbon which may be further functionalized by a member selected from the class consisting of amino, ether, epoxy, isocyanate, cyano, acryloxy and acyloxy and combinations thereof.

The C_{1-8} monovalent hydrocarbon radical of R^2 and R^3 includes an alkyl radical such as, for example, methyl,

ethyl, propyl, butyl, or an alkylene radical such as vinyl and allyl. Preferably R^2 and R^3 are alkyl radicals, more preferably methyl, ethyl or propyl radicals.

The saturated, unsaturated or aromatic hydrocarbon of Z includes alkyl, cycloalkyl, alkylene, and 5 aryl groups. Among the alkyl groups are C_{1-8} straight or branched-chain alkyl such as, for example, methyl, ethyl, propyl, isopropyl, n-butyl, isobutyl, tert-butyl, pentyl, isopentyl, hexyl, etc., the cycloalkyl are C_{3-8} cycloalkyl such as, for example, cyclopropyl, cyclobutyl, cyclohexyl, 10 etc., the alkylene groups are C_{1-8} alkylene such as, for example, vinyl, allyl, and the aryl groups include phenyl and napthyl. The above groups may be further functionalized by including in the chain or ring structure, as the case may be, a group selected from the class 15 consisting of amino, ether, epoxy, isocyanate, cyano, acryloxy, acyloxy and combinations. Preferably Z is an alkyl group, more preferably further functionalized by one or more amino group. The most preferred organo-functional silane is N-(2-aminoethyl-3-aminopropyl)trimethoxysilane. 20

The composition additionally contains about 0.05 to 1 weight percent of an organotin salt. Preferably the organotin salt is selected from the group dibutyltin diacetate, stannous octoate, dibutyltin dioctoate and dibutyltin dilaurate. Most preferably the organotin salt is dibutyltin dilaurate of the formula

 $(C_4H_9)_2Sn(OCOC_{10}H_{20}CH_3)_2$

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The composition also contains about 10 to 40 weight percent of a hydrocarbon solvent to be used as a dispersion medium for the composition. The hydrocarbon solvent is preferably a petroleum based solvent such as naphtha or mineral spirits.

The composition may contain other optional ingredients such as pigments and other fillers in minor

amounts provided that the addition of the ingredients does not cause degradation of the corrosion resistance of the cured coating made from the composition. One commonly utilized optional ingredient is a pigment, preferably a grey pigment, most preferably present in amounts up to about 1 weight percent.

The organopolysiloxane composition of the present invention is prepared by mixing the ingredients together in the absence of moisture. The silane is moisture sensitive and will undergo cross-linking in the presence of moisture such that the mixture must be essentially absent of free moisture when the silane is added and maintained in a moisture free state until cure is desired.

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A preferred method of mixing comprises mixing the polysiloxane polymer with the reinforcing fillers and other optional fillers and pigments. Thereafter, the oximinosilane and organo-gunctional silane are added and mixed under a nitrogen atmosphere. The solvent is added to the mixture under a nitrogen atmosphere and finally, the organotin salt is added to the mixture. The mixture is then dispensed in the sealed containers for storage prior to use.

The surface of the metal or concrete to be protected is coated with the composition by conventional methods such as dipping, brushing or spraying. Preferably, the metal or concrete to be protected is coated by spraying one or more applications of the composition of the present invention. The coating generally has an average thickness of 0.25 to 1.50 mm, single or more preferably, an average thickness of 0.5 to 1.0 mm, most preferably about 0.5 to 0.75 mm. After the coating is formed on the surface of the metal or concrete, the metal or concrete is exposed to normal atmosphere for cross-linking and cure of the coating.

The improved coating of the present invention is capable of protecting metal and concrete surfaces from corrosion in the presence of moisture such as rain or fog in combination with contaminated atmospheres, salt spray or fog or direct exposure to salt water.

The improved coating of the present invention is particularly useful for protecting metal surfaces which are directly exposed to salt water. Such surfaces include the hulls of ships and other vessels, oil drilling rigs, harbour and pier structures, etc. When the coating is used 10 on the hulls of ships, further benefits such anti-fouling in addition to the corrosion protection are achieved. coating does not allow marine animals, such as barnacles, to easily attach to the surface. Any such animals which attempt to attach to the surface are generally removed from 15 the surface by high pressure washers. Additionally, clean up of the surface is generally accomplished by high pressure wash and/or hand or mechanical wiping and does not require the scraping operations commonly utilized during hull cleaning of ships, or other marine installations. 20

The following examples are included to illustrate preferred embodiments of the invention and to demonstrate the usefulness of the coating and are not intended to limit in any way the scope of protection for the invention.

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EXAMPLE 1

To 36 parts of dimethyl polysiloxane fluid having a viscosity of 16.75 Pa.S at 25°C is added 35 parts of a mixture of anorphorous and crystalline silica fillers having a specific gravity of 2.2 and surface area of about $130\text{m}^2/\text{g}$. Then 2 parts of pigment is added and the composition is mixed in a mixer to a uniform consistency. Then 3 parts of methyl tris-(methyl ethyl ketoxime) silane and 1 part of N-(2-aminoethyl-3 aminopropyl)

trimethoxysilane are added and mixed under a nitrogen atmosphere. Then 22 parts of naphtha solvent is added to the mixture. Finally, 0.1 part of dibutyltin dilaurate is added to the dispersion and mixed until a uniform consistency is achieved.

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Coupon samples prepared from stainless and carbon steel and aluminum sheet were coated with the composition prepared according to Example 1 by dipping into the mixture and exposing the samples to normal atmosphere for crosslinking. The coupon samples, having an average coating thickness of 0.3 mm, were exposed to salt-fog testing in an electrical (ozone producing) environment. After 2000 hours of continuous exposure, the coating showed no evidence of deterioration or separation from the virgin metal surfaces. Uncoated samples of both metals showed severe etching and corrosion.

Carbon steel sheet samples were coated with the composition according to Example 1 by dipping into the mixture and exposing the samples to normal atmosphere for crosslinking. The samples, having an average coating thickness of 0.3 mm, were exposed to accelerated heat aging at 260°C. After 7 days of continuous exposure, the coating showed no evidence of deterioration or separation from the virgin metal surface.

25 Carbon steel coupons coated with the composition of Example 1 to a thickness of 0.5 mm were immersed 30 days in an algae producing environment of ocean water at room temperature. Considerable deposits of algae were present which were easily removed by wiping with a damp cloth. The original surface of the coating showed no changes, scratches or deformations.

The improved flexibility of the coating and its resistance to cracking was confirmed utilizing carbon steel coupons coated with 0.3 mm thick coating. The samples were

bent 180° along a curve showed no cracks or separation from substriate.

Compositions of the present invention are also useful for protecting other types of surfaces from corrosive environments. For example, compositions of the 5 present invention are particularly useful for protecting surfaces for spillage containment around tank farms and the like. The composition may be coated onto a heavy-duty fabric which is then utilized to line the interior of the spillage catch basins around the tank farm. Samples of 10 heavy-duty geo fabric were sprayed with the composition of the present invention to coat the fabric. Treated and untreated samples of fabric were exposed to caustic soda solution, diesel oil, furnace oil by placing samples of these materials on the surface of the fabric for up to 15 seven days. No deterioration of the treated fabric was observed and the deposits of the material were easily able to be cleaned up without any noticeable loss of weight of the material. In contrast, untreated fabric did not retain the material on the surface and the caustic soda solution 20 caused deterioration of the untreated fabric material within 24 hours.

While the invention has been described in
reference to specific embodiments it should be understood
by those skilled in the art that various changes can be
made and equivalents may be substituted without departing
from the true spirit and scope of the invention. All such
modifications are intended to be within the scope of the
claims appended hereto.

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE AS FOLLOWS:

- 1. A one-part organopolysiloxane rubber composition for use as a corrosion protection coating on metals comprising the product which is obtained by mixing the following:
 - a) about 20 to 50 weight percent of polydimethyl siloxane fluid of the formula

$HO[(CH_3)_2SiO]_nH$

in which n has an average value such that the viscosity is in the range from 1 to 100 Pa.S at 25°C, preferably from 10 to 20 Pa.S at 25°C;

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- b) about 20 to 50 weight percent of a mixture of anorphorous and crystalline SiO₂ reinforcing fillers having a surface area of up to 200 m²/g and a specific gravity of 2.2;
 - about 2 to 6 weight percent of an oximinosilane;
 - d) about 0.25 to 2 weight percent of an organofunctional silane;
- e) about 0.05 to 1 weight percent of an organotin salt; and
 - f) about 10 to 40 weight percent of an organic solvent as a dispersion medium for the above composition.
- 25 2. A composition according to claim 1 wherein the oximinosilane is a compound of the formula

RaSi(ON=CR'2)4-a

30 wherein R and R' represent a group of monovalent hydrocarbon radicals and the integer <u>a</u> has a value 1,

3. A composition according to claim 2 wherein R and R' are independently selected from the group consisting of alkyl and alkylene.

- 4. A composition according to claim 2 wherein R and S R' are each alkyl.
 - 5. A composition according to claim 2 wherein R and R' are independently selected from the group consisting of methyl, ethyl, propyl, butyl, vinyl and allyl.
- 6. A composition according to claim 4 wherein R and R' are independently selected from the group consisting of methyl and ethyl.
 - 7. A composition according to claim 1 wherein the organo-functional silane is a compound of the formula

15 R^{3}_{b} | ($R^{2}0$) 3-b - Si - 7

wherein R² and R³ are C₁₋₈ monovalent hydrocarbon 20 radicals, b varies from 0 to 3, preferably 0, Z is a saturated, unsaturated or aromatic hydrocarbon which may be further functionalized by a member selected from the class consisting of amino, ether, epoxy, isocyanate, cyano, acryloxy 25 and acyloxy and combinations thereof;

8. A composition according to claim 1 wherein the organo functional silane is a compound of the formula

NHCH2CH2NH2

CH₂

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CH₂CH₂Si (OMe)₃

wherein Me is the methyl radical.

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9. A composition according to claim 1 wherein the organotin salt is an organotin salt of a carboxylic acid selected from the group consisting of dibutyltindiacetate, stannous octoate and dibutyltin dioctoate.

10. A composition according to claim 9 wherein the organotin salt of a carboxylic acid is a compound of the formula

$(C_4H_9)_2Sn(OCOC_{10}H_{20}CH_3)_2$.

- 10 11. A composition according to claim 1 comprising:
 - a) about 35 weight percent of a hydroxyl terminated dimethyl polysiloxane fluid having a viscosity of 10 Pa.S at 25°C;
- b) about 35 weight percent of a mixture of
 anorphorous and crystalline SiO₂ fillers having a
 specific gravity of 2.2 and surface area of up to
 about 130m²/g;
 - c) about 4.3 weight percent of methyl tris-(methyl ethyl ketoxime)silane;
- 20 d) about 2.2. weight percent of N-(2 aminoethyl-3 aminopropyl) trimethoxysilane;
 - e) about 0.8 weight percent of dibutyltindilaurate;
 - f) about 22 weight percent of petroleum based solvent; and
- 25 g) about 0.8 weight percent of a grey pigment.
 - 12. A method of protecting a surface from the effects of a corrosive atmospheric environment comprising

(1) applying to the surface a thin layer of a onepart organopolysiloxane rubber composition comprising the product which is obtained by mixing the following:

a) about 20 to 50 weight percent of polydimethyl siloxane fluid of the formula

HO[(CH₃)₂SiO]_nH

in which n has an average value such that the viscosity is in the range from 1 to 100 Pa.S at 25°C, preferably from 10 to 20 Pa.S at 25°C;

- b) about 20 to 50 weight percent of a combination of anorphorous and crystalline SiO₂ reinforcing fillers having a surface area of up to 200 m²/g and specific gravity of 2.2;
 - c) about 2 to 6 weight percent of an oximinosilane;
- d) about 0.25 to 2 weight percent of an organo functional silane;
 - e) about 0.05 to 1 weight percent of an organtin salt; and
- f) about 10 to 40 weight percent of an organic solvent as a dispersion medium for the above composition.

and

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- (2) allowing the layer of the one-partorganopolysiloxane rubber composition to cure at roomtemperature to a silicone elastomer.
 - 13. A method according to claim 12 wherein the one part organopolysiloxane rubber composition comprises:
- a) about 35 weight percent of a hydroxyl terminated dimethyl polysiloxane fluid having a viscosity of 10 Pa.S at 25°C;

b) about 35 weight percent of a mixture of anorphorous and crystalline SiO_2 fillers having a specific gravity of 2.2 and surface area of up to about $130m^2/g$;

- 5 c) about 4.3 weight percent of methyl tris-(methyl ethyl ketoxime) silane;
 - d) about 2.2. weight percent of N-(2 aminoethyl-3 aminopropyl) trimethoxysilane;
 - e) about 0.8 weight percent of dibutyltindilaurate;
- 10 f) about 22 weight percent of petroleum based solvent; and
 - g) about 0.8 weight percent of a grey pigment.
- 14. A method according to Claims 12 or 13 wherein the surface to be protected is a metal or concrete surface exposed to a corrosive salt water environment.

INTERNATIONAL SEARCH REPORT

Im sional Application No PCT/CA 98/00673

A. CLASSI IPC 6	FICATION OF SUBJECT MATTER C09D5/08 C08K5/54 C08K3/3	6 C09D183/04		
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Electronic d	tata base consulted during the international search (name of data ba	ase and, where practical, search terms used)	
C. DOCUM	ENTS CONSIDERED TO BE RELEVANT			
Category *	Citation of document, with indication, where appropriate, of the re	levant passages	Relevant to claim No.	
Y	EP 0 340 120 A (RHONE-POULENC) 2 November 1989 see page 8, line 31 - page 9, li	ne 7·	1-14	
	claims 1,5	,,		
Y	EP 0 143 541 A (TORAY) 5 June 198 see page 7, line 20 - line 24 see page 9, line 15 - line 25	85	1-14	
Y	US 4 931 491 A (SAVIN) 5 June 19 see column 4, line 22 - line 36	90	1-14	
A	EP 0 735 101 A (DOW CORNING TORA) 2 October 1996 see claim 1; example 1	Y)	1-14	
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X Furt	her documents are listed in the continuation of box C.	X Patent family members are listed	in annex.	
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Int. Ional Application No PCT/CA 98/00673

	tion) DOCUMENTS CONSIDERED TO BE RELEVANT	
egory *	Citation of document, with indication,where appropriate, of the relevant passages	Relevant to claim No.
	US 4 734 479 A (INOUE ET AL) 29 March 1988 see column 7, line 60 - column 8, line 11; claim 1	1-14
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INTERNATIONAL SEARCH REPORT

information on patent family members

Inti Ional Application No PCT/CA 98/00673

	itent document I in search repor	t	Publication date	ı	Patent family member(s)	Publication date
EP	340120	A	02-11-1989	FR	2629827 A	13-10-1989
				AU	3251889 A	12-10-1989
	•			CA	1316286 A	13-04-1993
				JP	1306470 A	11-12-1989
				US	4996112 A	26-02-1991
EP	143541	Α	05-06-1985	JP	60096650 A	30-05-1985
				AU	575868 B	11-08-1988
				AU	3484284 A	09-05-1985
				BR	8405243 A	27-08-1985
				CA -	1247775 A	27-12-1988
				US	4535109 A	13-08-1985
US	4931491	Α	05-06-1990	NONE		
EP	735101	A	02-10-1996	JP	8269335 A	15-10-1996
				CA	2172915 A	01-10-1996
				US	5780543 A	14-07-1998
US	4734479	A	29-03-1988	JP	1669993 C	12-06-1992
				JP	3033749 B	20-05-1991
				JP	62223264 A	01-10-1987